# **DEPARTMENT OF JUSTICE**

#### **Drug Enforcement Administration**

# Importation of Controlled Substances; Notice of Application

Purusant to Section 1008 of the Controlled Substances Import and Export Act (21 U.S.C. 958(i)), the Attorney General shall, prior to issuing a registration under this Section to a bulk manufacturer of a controlled substance in Schedule I or II and prior to issuing a regulation under Section 1002(a) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with Section 1301.34 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on March 19, 1999, Noramco of Delaware, Inc., Division of McNeilab, Inc., 500 Old Swedes Landing Road, Wilmington, Delaware 19801, made application by renewal to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Opium, raw (9600)	II
Opium granulated (9640)	II

The firm plans to import the listed controlled substances for the bulk manufacture of other controlled substances.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of these basic classes of controlled substances may file written comments on or objections to the application described above and may, at the same time, file a written request for a hearing on such application in accordance with 21 CFR 1301.43 in such form as prescribed by 21 CFR 1316.47.

Any such comments, objections or requests for a hearing may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than June 7, 1999.

This procedure is to be conducted simultaneously with and independent of the procedures described in 21 CFR 1301.34(b), (c), (d), (e), and (f). As noted in a previous notice at 40 FR 43745–46 (September 23, 1975), all applicants for registration to import a basic class of

any controlled substance in Schedule I or II are and will continue to be required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration that the requirements for such registration pursuant to 21 U.S.C. 958(a), 21 U.S.C. 823(a), and 21 CFR 1301.34(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: April 26, 1999.

#### John H. King,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 99–11450 Filed 5–6–99; 8:45 am] BILLING CODE 4410–09–M

#### **DEPARTMENT OF JUSTICE**

### **Drug Enforcement Administration**

# Manufacturer of Controlled Substances; Notice of Registration

By Notice dated January 27, 1999, and published in the **Federal Register** on February 4, 1999, (64 FR 5689), Orpharm, Inc., 4815 Dacoma, Houston, Texas 77072, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Methadone (9250)	    

The firm plans to manufacture methadone and methadone-intermediate for production of LAAM.

DEA has considered the factors in 21 U.S.C. § 823(a) and determined that the registration of Orpharm, Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Orpharm, Inc. on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the company's physical security systems, audits of the company's records, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to 21 U.S.C. 823 and 28 C.F.R. 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic classes of controlled substances listed above is granted.

Dated: April 26, 1999.

#### John H. King,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

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#### **DEPARTMENT OF JUSTICE**

# Immigration and Naturalization Service

[INS No. 1990-99]

# Announcement of District Advisory Council on Immigration Matters Sixth Meeting

**AGENCY: Immigration and Naturalization** 

Service, Justice.

**ACTION:** Notice of meeting.

SUMMARY: The Immigration and Naturalization Service (Service) has established a District Advisory Council on Immigration Matters (DACOIM) to provide the New York District Director of the Service with recommendations on ways to improve the response and reaction to customers in the local jurisdiction, and to develop new partnerships with local officials and community organizations to build and enhance a broader understanding of immigration policies and practices. The purpose of this notice is to announce the forthcoming meeting.

DATES AND TIMES: The Sixth meeting of the DACOIM is scheduled for May 27, 1999, at 1 p.m.

ADDRESSES: The meeting will be held at Orange County Community College, 115 South Street, Middletown, New York, 10940, in the Biotech Building (between Ramview and South Street), Room 207.

FOR FURTHER INFORMATION CONTACT: Susan Young, Designated Federal Officer, Immigration and Naturalization Service, 26 Federal Plaza, Room 14–100, New York, New York, 10278, telephone: (212) 264–0736.

**SUPPLEMENTARY INFORMATION:** Meetings will be held tri-annually on the fourth Thursday during the months of January, May, and September 1999.

# **Summary of Agenda**

The purpose of the meeting will be to conduct general business, review subcommittee reports, and facilitate public participation. The DACOIM will be chaired by Charles Troy, Assistant District Director for Management, New York District, Immigration and Naturalization Service.

#### **Public Participation**

The DACOIM meeting is open to the public, but advance notice of attendance